

Rediscovery of the endemic gastropod *Dianella schlickumi* (Gastropoda, Hydrobiidae) and its discrimination from *Dianella thiesseana*: environmental correlates and implications for their conservation

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Academic editor: M. Pinna | Received 7 January 2018 | Accepted 24 May 2018 | Published 11 June 2018

<http://zoobank.org/EF862219-884C-436F-BF54-7603F97EEFFB>

Citation: Ntislidou C, Radea C, Giokas S, Pusch MT, Lazaridou M, Bobori DC (2018) Rediscovery of the endemic gastropod *Dianella schlickumi* (Gastropoda, Hydrobiidae) and its discrimination from *Dianella thiesseana*: environmental correlates and implications for their conservation. Nature Conservation 27: 35–58. <https://doi.org/10.3897/natureconservation.27.23289>

Abstract

The aquatic snail genus *Dianella* (Gastropoda: Hydrobiidae) has only two representatives in Greece: *Dianella schlickumi* Schütt, 1962 and *Dianella thiesseana* (Kobelt, 1878). *D. schlickumi*, a narrow endemic species to Lake Amvrakia (in Aitolioakarnania, western-central Greece), is considered as Critically Endangered (Possibly Extinct, sensu IUCN 2017). Our study confirmed its presence in Lake Amvrakia, where it had not been detected for more than 30 years. We document the unknown anatomical characters based on the *D. schlickumi* specimens. Moreover, the presence of *D. thiesseana* in the nearby lakes Trichonis and Lysimachia was also confirmed, while morphometric analyses enabled the discrimination between the two species. Redundancy Analysis revealed conductivity, dissolved oxygen and pH as the main environmental variables related to the above species' distribution, shaping their community structure. Both *Dianella* species require urgent conservation measures to be enforced, due to their habitat degradation from human activities, which are limiting and fragmenting their range. For that purpose, effective management plans have to be elaborated and implemented at the mentioned lakes, focusing on the reduction of human pressures and on the improvement of their habitats.

Keywords

Pyrgulinid, Greek lakes, anatomy, morphometrics, environmental correlates, conservation

Introduction

Freshwater habitats cover less than 1% of the earth's surface, but they support almost 10% of the known species on the planet (Strayer and Dudgeon 2010). However, freshwater species are in a worldwide decline (Howard et al. 2015), resulting thus in the decrease of freshwater biodiversity, which is mainly threatened by over-exploitation, water pollution, hydrological alterations, destruction and degradation of habitats and invasion by exotic species (Dudgeon et al. 2006).

The Balkan region is rich in freshwater fauna due to high endemism (Griffith et al. 2004, Glöer et al. 2007). The rich fauna of freshwater molluscs is considerably threatened (Albrecht and Wilke 2008) and 29 species of the Balkan Peninsula have already become extinct (Régner et al. 2009), representing 21% of the 140 known extinctions of freshwater mollusc species worldwide. In this region, many endemic gastropod species exist with a restricted range only extending to small hydrographic systems (rivers, lakes and springs). The karst landscape, which occurs in a large proportion of the Balkan Peninsula, also contributes to this high degree of endemism (Régner et al. 2009). These species are vulnerable not only due to habitat degradation caused by human activities, but also by their very limited range, which increases the risk of accidental extinction (Régner et al. 2009). For Greece, a similar crucial decline of population densities and losses of some endemic mollusc species have been reported (Albrecht et al. 2006), suggesting the necessity for urgent actions for their conservation.

The distribution of gastropod species is often associated with biotic and abiotic habitat parameters (e.g. Covich 2010), which are related to specific geographical characteristics, physicochemical conditions, biological interactions and historical and random factors (Lodge et al. 1987). Thus, the distributional patterns of endemics can be often better explained based on records of key environmental parameters, which drive the structure of biological communities (Pérez-Quintero 2012). Such knowledge may also help to identify and protect specific regions for the conservation of biodiversity of little-known freshwater bodies (Abell et al. 2007), especially in the Mediterranean region which has been neglected.

The species of the subfamily Pyrgulinae (Caenogastropoda: Truncatelloidea: Hydrobiidae) are widely distributed in western Asia as well as in central and south-eastern Europe (Wilke et al. 2007). In the latter area, the subfamily consists of ten genera, namely *Chilopyrgula* Brusina, 1896, *Dianella* Gude, 1913, *Ginaia* Brusina, 1896, *Neofossarulus* Polinski, 1929, *Ochridopyrgula* Radoman, 1955, *Pyrgohydrobia* Radoman, 1955, *Pyrgula* De Cristofori & Jan, 1832, *Stankovicia* Polinski 1929, *Trachyochridia* Polinski, 1929 and *Xestopyrgula* Polinski, 1929, comprising 17 species (Bank 2017).

The pyrgulinids are mostly lacustrine snails, colonising the littoral and sublittoral zones of ancient deep freshwater lakes (Radoman 1983, Wilke et al. 2007, Schreiber et al. 2012). Only one species, *Chilopyrgula sturanyi* Brusina, 1896, has been recorded in springs (Radoman 1983).

Out of those, two pyrgulinid species, belonging to the genus *Dianella* Gude, 1913, are known to be present in lentic systems of Greece: *Dianella schlickumi* Schütt, 1962 and *Dianella thiesseana* (Kobelt, 1878). The first species is known from Lake Amvrakia and the latter from lakes Trichonis and Lysimachia (Aitolioakarnania, western-central Greece) (Schütt 1962, Radoman 1973, 1983, Szarowska et al. 2005, Szarowska 2006). Both species are narrow endemic species, with major gaps in historical records, while no information is available about their abundance, population structure, life cycle or genetic diversity. In the IUCN Red List of Threatened Species, *Dianella thiesseana* appears under the status “Critically Endangered”, while *D. schlickumi* has been declared as “Possibly Extinct” (IUCN 2017). Apart from *Dianella*, the aquatic gastropod communities of these lakes include several other endemic hydrobiids, such as *Islamia graeca* Radoman, 1973, *Islamia trichoniana* Radoman, 1979, *Pseudoislamia balcanica* Radoman, 1979, *Trichonia trichonica* Radoman, 1973 (Radoman 1983). *D. schlickumi* has not been observed for more than 30 years, despite several sampling efforts performed in Lake Amvrakia (Reischütz and Reischütz 2002, Szarowska et al. 2005, Albrecht et al. 2006, 2011a), which is considered as the type locality of this species. Only fresh empty shells of this species were recorded in an irrigation channel, near Acheloos River (close to Lake Amvrakia) by Reischütz et al. (2008). *D. thiesseana* has only been found at the northeast bank of Lake Trichonis during the last decade (Albrecht et al. 2009, 2011b).

The external morphology, the anatomy and the phylogenetic position of the majority of Pyrgulinae have been extensively studied and discussed (Schütt 1962, Radoman 1983, Riedel et al. 2001, Szarowska et al. 2005, Szarowska 2006, Anisratenko 2008, Wilke et al. 2013). The combination of anatomical and phylogenetic data suggests that the Pyrgulinae subfamily has a very close (possibly even sister-group) systematic relationship with Hydrobiinae (Szarowska et al. 2005). Concerning the two species of *Dianella*, it has been reported that they have similar external morphology and soft body anatomy, but different body size (Radoman 1983). However, the inter-specific differences have been questioned by Szarowska et al. (2005), which has made the taxonomic distinction between *D. thiesseana* and *D. schlickumi* unclear and questionable. Although the soft body anatomy of *D. thiesseana* is described and/or depicted in detail (Radoman 1983, Szarowska et al. 2005, Szarowska 2006), various anatomical characters of *D. schlickumi* remain unknown, un-described or unpublished.

Hence, the aims of the present study were to: (a) report the rediscovery of the seemingly extinct pyrgulinid *D. schlickumi*, (b) describe its unknown anatomical characters, (c) examine its morphometric discrimination from *D. thiesseana* and (d) reveal the environmental parameters that probably drive the distribution of both species.

Material and methods

Study area

We studied three natural lakes located in the western-central part of Greece (Figure 1). These lakes are considered as the last remains of a former big lake, which by several processes, has been split into four smaller lakes (Albanakis et al. 1995). The Acheloos River separates Amvrakia and Ozeros lakes from Trichonis and Lysimachia lakes (Figure 1). Lake Amvrakia (Table 1) is a semipolje karst lake of tectonic origin with high sulphate concentrations, situated in Mesozoic limestone (Verginis and Leontaris 1978, Overbeck et al. 1982). Its water level shows significant seasonal fluctuations, which may be attributed to various factors such as the hydraulic communication with karst aquifers, high evaporation rates, especially during summer and intensive use of water for agricultural purposes (Danielidis et al. 1996). Lake Trichonis (Table 1) is the deepest and the largest natural lake in Greece, which is also a karst water body (Zacharias et al. 2002), situated in a tectonically active area (Poulimenos and Doutsos 1997). It is supplied by precipitation, approximately 30 intermittent streams and sub-aquatic karst springs (Dimitriou and Zacharias 2006). At its western end, a narrow canal connects Lake Trichonis with Lake Lysimachia (Table 1), which is shallower (Figure 1) and it was receiving the untreated urban wastewaters from the city of Agrinio until the year 2000. At its western end, Lake Lysimachia drains to the Acheloos River via an artificial canal (Figure 1).

Field sampling

Macroinvertebrate samplings were conducted in spring and autumn 2014 by boat, at the sublittoral and profundal zones of each lake, using an Ekman-Birge grab (three replicates per station, 225 cm² sampling area). A total of 66 samples were collected in the studied lakes [Amvrakia: 14 stations (7 stations at each zone), Trichonis: 13 (5 and 7 stations at the sublittoral and profundal zones, respectively) and Lysimachia: 6 (3 stations at each zone); Figure 1]. Samples were sieved with a 500 µm mesh and preserved

Table 1. Geographical position, morphological and limnological features of the studied lakes. Alt: altitude; Z_{mean} : mean depth; Z_{max} : max depth; L_a : lake surface area; TL: trophic level, ET: eutrophic, MT: mesotrophic, OL: oligotrophic; WMM: warm monomictic.

Lake	Latitude	Longitude	Alt (m.a.s.l)	Z_{mean} (m)	Z_{max} (m)	L_a (km ²)	TL	Lake type
Trichonis	38.573333	21.552222	18 [†]	30 [†]	57 [‡]	96.5	OL/MT [§]	WMM [†]
Lysimachia	38.558597	21.376873	16 [†]	3 [‡]	9 [‡]	13.1	ET [‡]	WMM [‡]
Amvrakia	38.652150	21.219867	16 [†]	23	53 [¶]	11.8	MT [#]	WMM [#]

[†]Zacharias et al. (2002), [‡]Skoulidakis et al. (1998), [§]Doulka and Kehayias (2008), [‡]Leonardos (2004), [¶]Albanakis et al. (1995), [#]Danielidis et al. (1996)

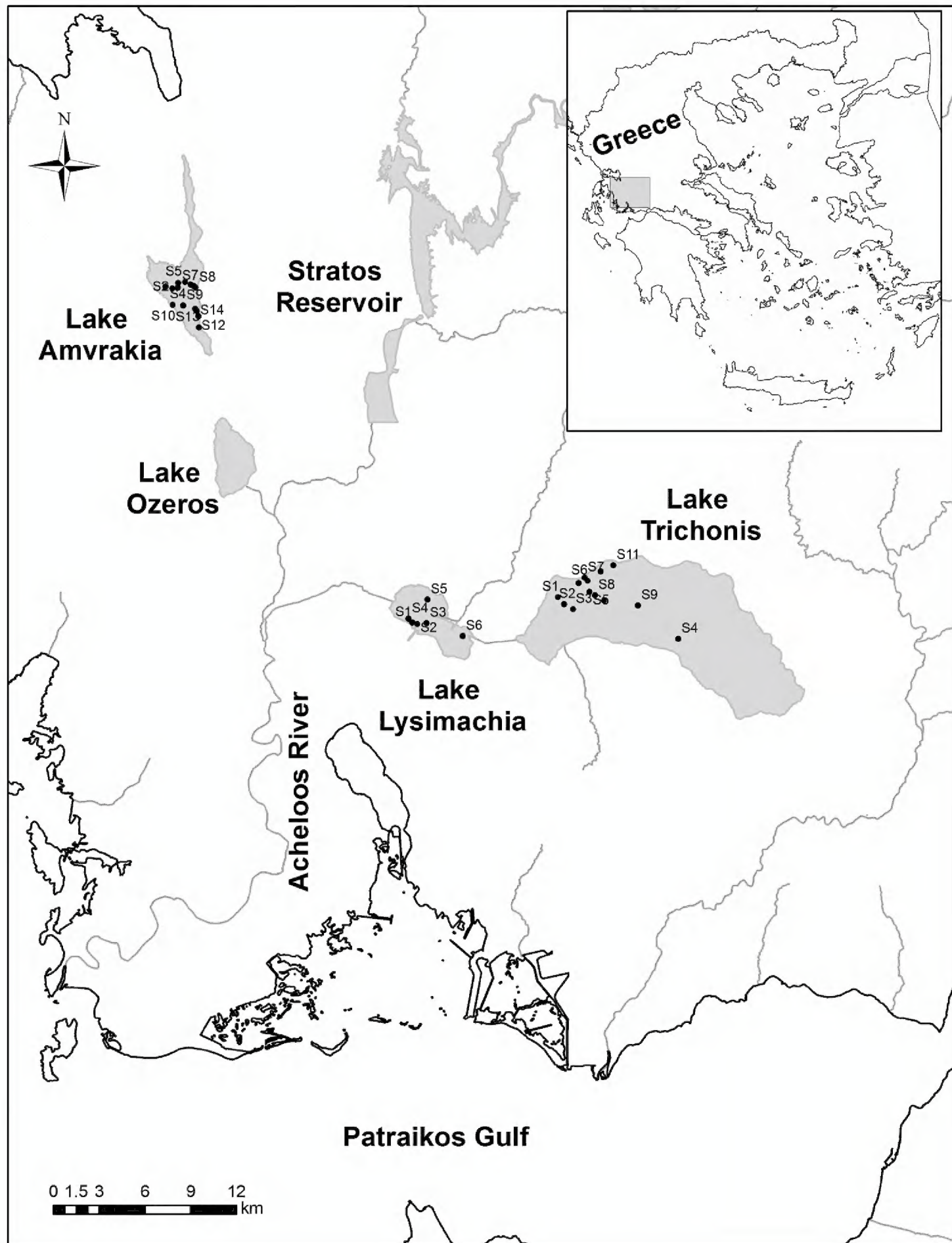


Figure 1. Geographical location of the studied lakes and sampling stations.

in 70% ethanol. After sorting, zoobenthos was identified to the lowest possible taxon and abundance was converted to density (ind./m²). Additionally, at each station, water samples from 1 m above the bottom of each lake were collected, using a Niskin-type sampler. Environmental parameters such as water temperature (WT, °C), pH, conduc-

tivity (Cond, $\mu\text{S}/\text{cm}$) and dissolved oxygen concentration (DO, mg/l) were measured *in situ* just above the bottom sediment of each station using the Aqua Read AP-2000 probe (Kent, GB). Secchi depth (Secchi, cm) was also recorded.

Anatomy

Genitalia of male and female specimens and other soft body characteristics of *D. schlickumi* from Lake Amvrakia were studied and compared with those of *D. thiesseana* collected from Lake Trichonis during the present study and with some additional individuals collected in 2016 (Figure 2). Before dissection, the shells of the specimens studied were removed by soaking in Perenyi solution, following Hershler and Ponder's (1998) morphological terminology. The characterisation of the nervous system was based on Davis et al. (1986) and its concentration was measured as the RPG ratio (Davis et al. 1976).

Morphometry

Shell morphometry was recorded in both *D. thiesseana* (Trichonis: 33 individuals and Lysimachia: 5 individuals) and *D. schlickumi* (Amvrakia: 44 individuals) in well preserved shells of adult specimens, using the software ImageFocus v3.0.0.1. Seven linear measurements were taken, specifically: shell height (H), shell width (W), shell aperture height (Ha), shell aperture width (Wa), spire height (SH), body whorl height (BWH) and penultimate whorl height (PWH) (Figure 3). Measurements were then expressed as ratios: shell height/shell width (H/W); shell aperture height/shell aperture width (Ha/Wa); spire height/body whorl height (SH/BWH), penultimate whorl height/body whorl height (PWH/BWH) and body whorl height/shell width (BWH/W). Prior to further analyses, all the linear measurements and the ratios were logarithmically transformed for normalisation. Subsequently, the t-test for independent samples was performed for detecting significant differences ($p < 0.05$) between the two species using the statistical software SPSS v22.

The studied populations were also tested for morphological diversification using a landmark-based method of acquiring geometrical data of shape and size. Shell variation has been traditionally quantified through linear measurements and ratios to distinguish between individuals and populations, amongst and within snail species. Recently, geometric morphometrics (GMs) have been employed for examining shells, both to provide direct size-free analyses of shell shape (Conde-Padin et al. 2007b, Hayes et al. 2007) and to answer broader evolutionary questions (Pfenninger and Magnin 2001, Conde-Padin et al. 2007a, Haase and Misof 2009, Páll-Gergely et al. 2012, Giokas et al. 2014). GMs have the advantage over traditional morphometrics in being, theoretically, free of effects due to size, position, rotation and scale (Rohlf and Marcus 1993).

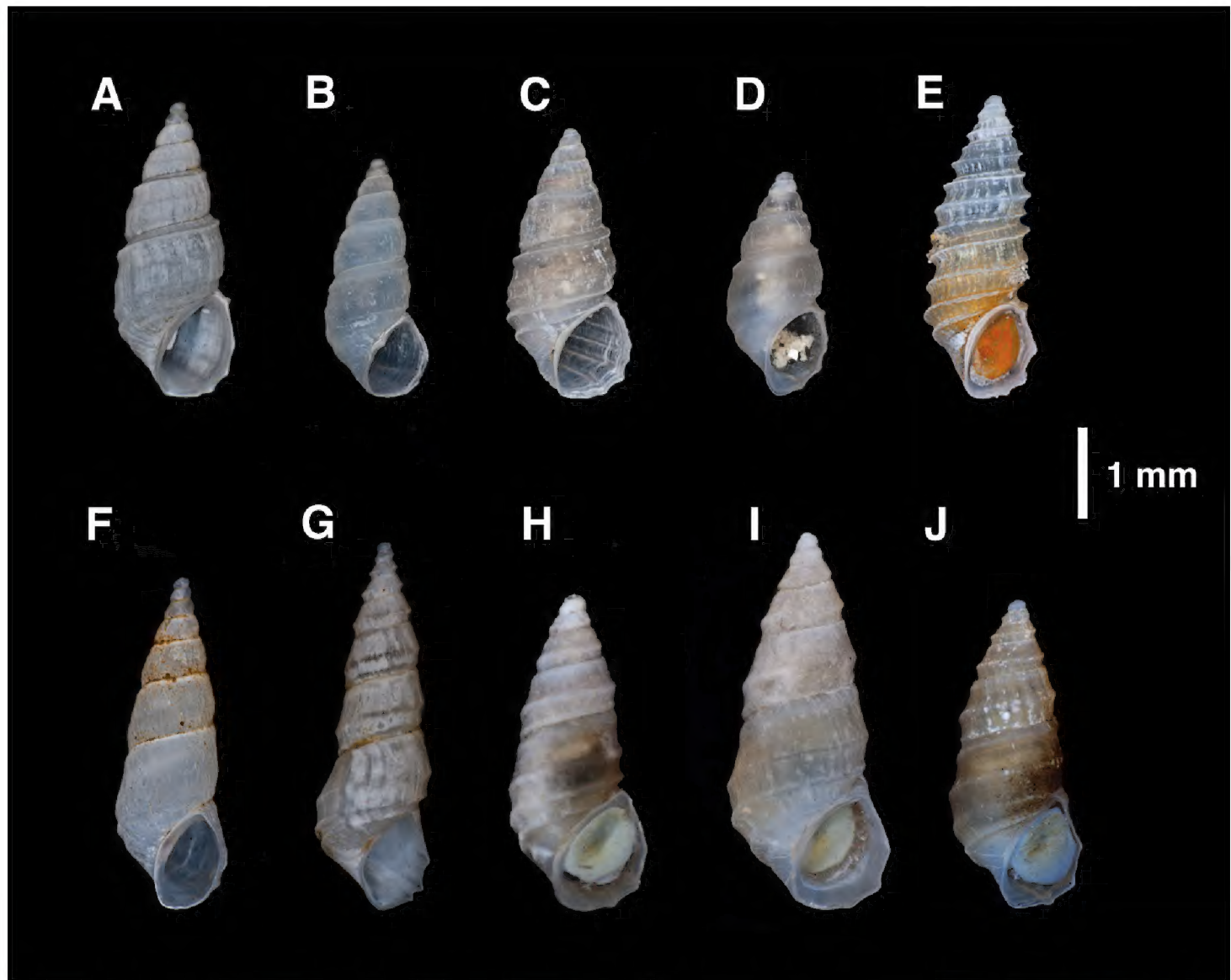


Figure 2. Specimens of: *Dianella schlickumi* from Lake Amvrakia (**A–E**) and *Dianella thiesseana* from Lake Trichonis (**F–J**), demonstrating their large inter-specific variation of shell morphology.

For that purpose, we used sub-samples of the undamaged and well preserved shells of adult specimens (i.e. 15 *D. schlickumi* and 9 *D. thiesseana* specimens), using the aperture features. Specimens were set down on the same plane, with the aperture facing up and we took digital photographs of them. Geometric morphometric variables of the shells were obtained with 18 landmarks (LM) representing the outline of the shell and of the aperture as shown in Figure 4. Landmarks were digitised using tpsDig (Rohlf 2006) and occupied the same positions over the sum of the specimens. Two of these (LM1 and LM18) were of type I (developmentally homologous), 12 landmarks (LM2-LM13) represented suture lines and were considered type II (geometrically homologous). The remaining landmarks were either extreme points on surfaces or were placed halfway between other landmarks, making them type III (Bookstein 1991, Zelditch et al. 2004).

All geometric morphometric analyses were performed with MorphoJ (Klingenberg 2011). The coordinates of shape to be used for further statistical analysis were obtained with Procrustes generalised least squares superimposition. This method excludes the impact of size on the shape of the shell and, theoretically, variations independent of shape are removed using this analysis (Zelditch et al. 2004). Size variables

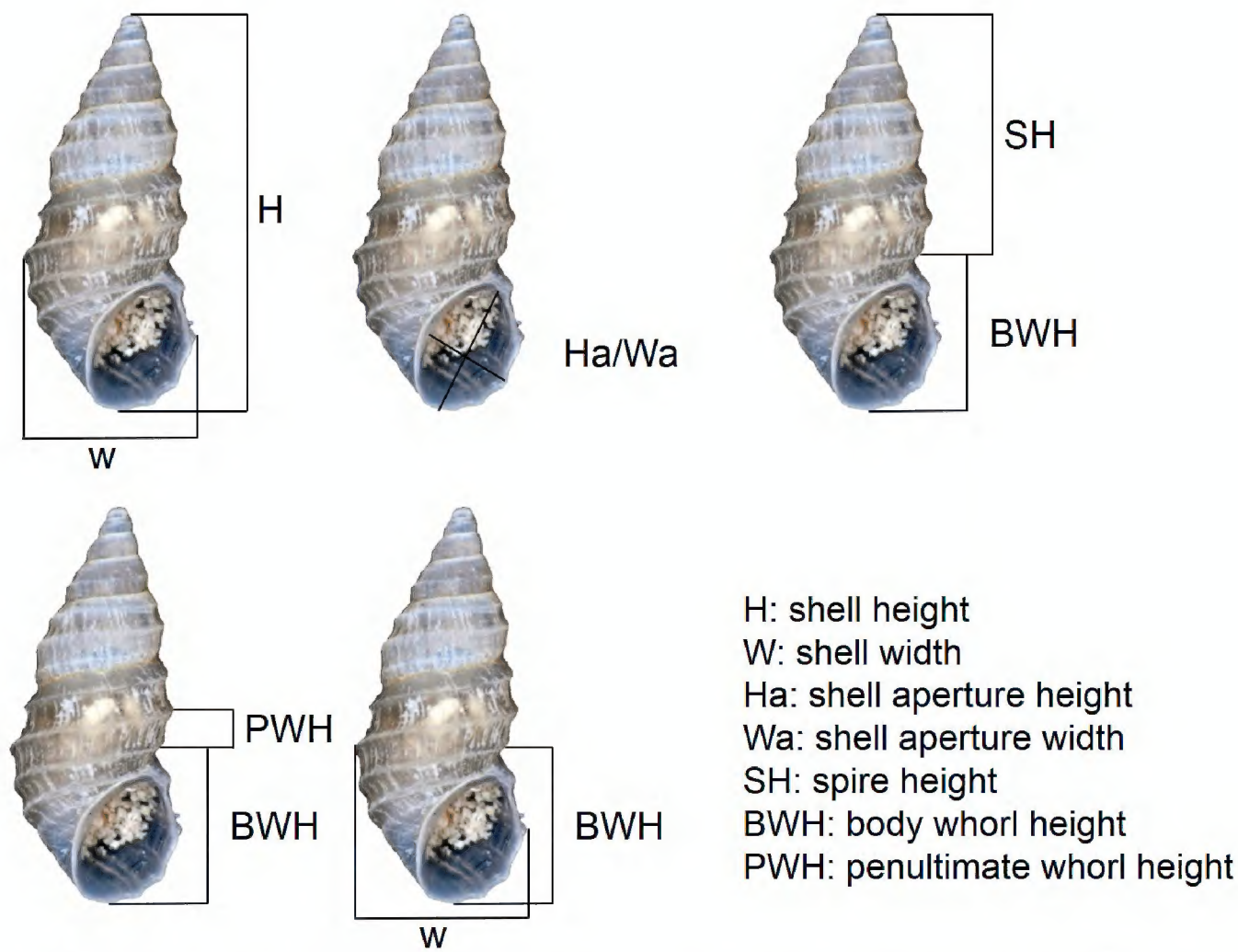


Figure 3. External morphological measurements taken in specimens of *Dianella schlickumi* and *Dianella thiesseana*.

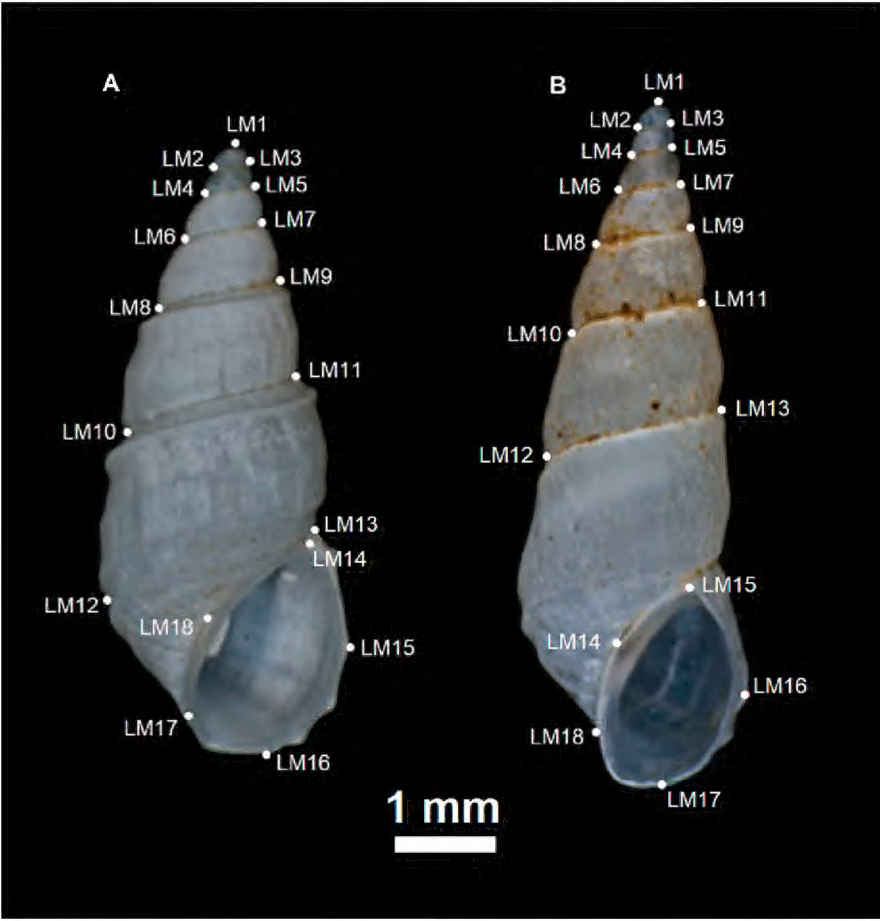


Figure 4. *Dianella schlickumi* and *Dianella thiesseana* specimens positioned with the axis of the shell on the y-axis and the aperture in the same plane as the objective, showing the location of the 18 landmarks (LM1-LM18) used. **A** *D. schlickumi* **B** *D. thiesseana*.

(centroid size) for each specimen were also generated. We examined, using pooled within-group regression analysis, the effect of centroid size on shape, regressing to the resulting Procrustes coordinates on centroid size. That regression score was not significant (Permutation test, 10,000 rounds against the null hypothesis of independence, % predicted = 6.170, $p = 0.2183$). Therefore, we eventually used the Procrustes coordinates, which were analysed with Principal Component Analysis (PCA) and Canonical Variate Analysis (CVA) in order to explore the morphological variation between the two *Dianella* species. The reliability of the discrimination was assessed by leave-one-out cross-validation.

Analyses of abundance and environmental data

Detrended Correspondence Analysis (DCA) (indirect gradient analysis) (programme CANOCO version 4.5.1; Ter Braak and Šmilauer 1998) was performed to abundance data in order to verify if they corresponded linearly to the environmental gradient or if abundances peaked around an environmental optimum (unimodal response) (Ter Braak and Šmilauer 1998). In this study, the species abundance data corresponded roughly linearly to the gradient, because the length of the gradient of the first axis was three times less than the range of the within-sample standard deviation (Ter Braak and Šmilauer 1998). Thus, Redundancy Analysis (RDA) was further applied to link the environmental parameters to species abundance data. The Monte Carlo permutation test was performed to select the statistically significant ($p < 0.05$) environmental parameters that explain most of the variance of species abundance, while the variation inflation factor (>20) was used to assess the interrelation of the parameters.

Results

Species distribution

Dianella schlickumi was found at the south-east part of Lake Amvrakia, in 5 (S7, S11, S12, S13, S14; Figure 1) out of the 14 stations (36%) surveyed, at depths ranging from 5 m to 13 m. No individuals were recorded at its western part (S9 and S10, Figure 1), not even empty shells. The highest abundance was recorded at station S15 (607 ind./m²) at depth 5 m (Table 2). *D. thiesseana* was found in 2 (S6 and S13) out of the 13 stations (15%) in Lake Trichonis, located on the northeast side of the lake, at depths of 13 m to 18 m (Figure 1, Table 2). During the current study, only empty shells from *D. thiesseana* were collected in Lake Lysimachia. However, the presence of a live *D. thiesseana* was confirmed in the east side of Lake Lysimachia by surveys executed during the National Water Monitoring Programme in spring 2015 (Mavromati, personal communication). Both species were present during the spring surveys while none was found during autumn.

Table 2. *Dianella schlickumi* and *Dianella thiesseana* abundances (number of individuals sampled by Ekman-Birge grab, 225 cm² sampling area) per sampling station and environmental parameters measured in lakes Trichonis, Lysimachia and Amvrakia.

Lake	Station	Abundance (ind./m ²)		Environmental parameters					
		<i>Dianella schlickumi</i>	<i>Dianella thiesseana</i>	Depth (m)	Secchi (m)	DO (mg/l)	pH	Cond (μS/cm)	WT (°C)
Trichonis	S6		59	18.0	6.4	9.05	10.04	319	18.8
	S13		44	13.0	5.6	8.92	10.73	320	15.0
Lysimachia	LYS		15	3.8	1.7	9.77	8.13	363	11.9
Amvrakia	S7	15		13.0	5.2	8.87	9.74	982	21.2
	S11	15		8.0	4.7	8.38	10.10	921	20.5
	S12	44		11.5	4.8	8.48	10.48	922	20.5
	S13	44		9.0	5.1	8.66	10.42	916	21.6
	S14	607		5.0	3.2	8.47	10.62	898	20.9

Soft body anatomy of *D. schlickumi*

Ctenidium-Osphradium: Ctenidial filaments broader than high; osphradium elongate, approximately opposite to the middle of ctenidium.

Nervous system (Figure 5A): Cerebral ganglia similarly sized, white; supraoesophageal and suboesophageal ganglia white; supraoesophageal connective much longer than the suboesophageal. Nervous system extremely elongate, RPG ratio 0.75.

Gastric caecum: Large and elongate gastric caecum on the posterior stomach chamber.

Rectum: The U-shaped intestine loop in the pallial cavity roof was wide and the faecal pellets were packed sideways (Figure 5B).

Female reproductive system (Figure 6A): Pallial oviduct glands with straight border; bursa copulatrix medium-sized, ovoid-globular, positioned posteriorly to the albumen gland; bursal duct long; renal oviduct unpigmented, coiled tightly with more than two bends; typical seminal receptacles absent; a beak-shaped dilatation of oviduct in the position of seminal receptacle rs2, anteriorly to the bursa.

Penis (Figure 6B): Penis large, gradually tapered, proximal part folded, penial base of medium width, its attachment area central and well behind the head, penial duct almost straight located at the outer edge of the penis.

Egg capsule (Figure 6C–D): Several lenticular and transparent egg capsules (each one containing a single egg) in various growth stages were found on the external shell surface of two specimens.

Morphometry

Seven out of the 12 morphometric measurements taken, differed statistically ($p < 0.001$) between the two species. Generally, individuals of *D. thiesseana* exhibited higher mean values compared to *D. schlickumi*, except for SH/BWH ratio, which was higher in *D. schlickumi* (Table 3).

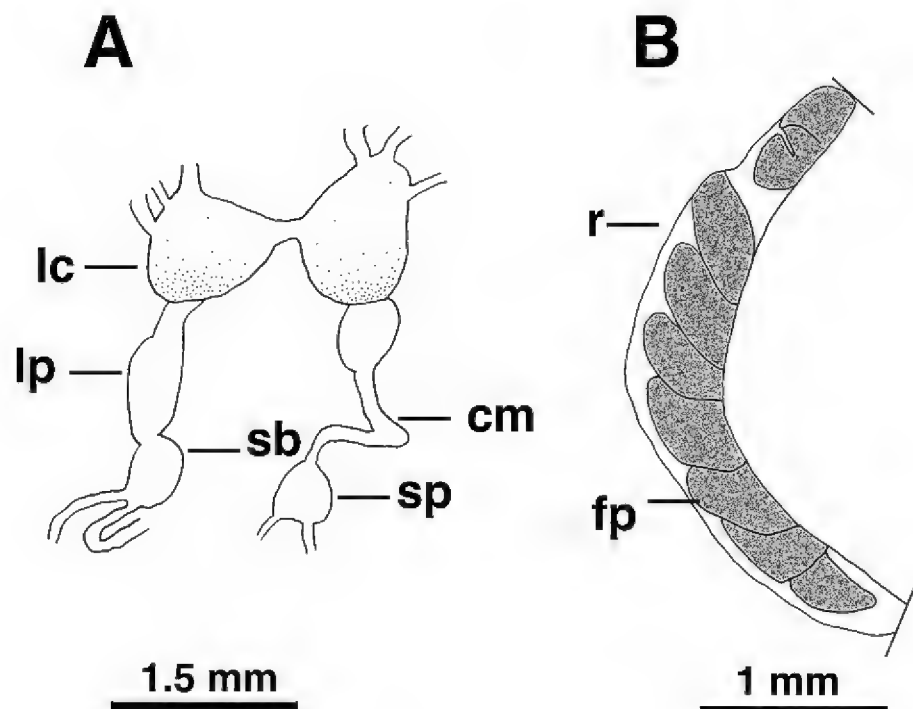


Figure 5. Non-genital anatomy of *Dianella schlickumi*. **A** Nervous system **B** Rectum full of faecal pellets. Abbreviations: cm -commissure; fp -faecal pellets; lc -left cerebral ganglion; lp -left pleural ganglion; sb - suboesophageal ganglion; sp -supraoesophageal ganglion; r -rectum.

Table 3. Average (\pm standard error) and range (minimum and maximum values) of the morphometric measurements (in mm) taken in *Dianella thiesseana* and *Dianella schlickumi* specimens. The results of t-test are also provided (* indicates the significant differences). n = number of individuals used: H: shell height; W: shell width; Ha: shell aperture height; Wa: shell aperture width; SH: spire height; BWH: body whorl height; PWH: penultimate whorl height.

Variables	<i>Dianella thiesseana</i> (n=38)	<i>Dianella schlickumi</i> (n=44)	t-test	
			<i>F</i>	<i>p</i>
H	6.79 \pm 0.25 (4.16–9.78)	6.17 \pm 0.21 (4.34–10.89)	3.95	0.063
W*	3.01 \pm 0.07 (2.20–3.86)	2.65 \pm 0.06 (1.93–3.75)	0.40	<0.001
Ha*	2.45 \pm 0.06 (1.58–3.16)	2.07 \pm 0.04 (1.61–2.92)	1.45	<0.001
Wa*	1.67 \pm 0.04 (1.18–2.22)	1.45 \pm 0.03 (1.05–1.91)	0.04	<0.001
SH	4.10 \pm 0.19 (2.05–6.38)	3.97 \pm 0.17 (2.50–7.70)	8.87	0.797
BWH*	2.69 \pm 0.07 (2.00–3.77)	2.20 \pm 0.05 (1.64–3.25)	0.86	<0.001
PWH*	0.67 \pm 0.02 (0.34–0.92)	0.54 \pm 0.01 (0.34–0.78)	5.23	<0.001
H/W	2.24 \pm 0.06 (1.69–3.14)	2.31 \pm 0.04 (1.99–3.29)	3.09	0.250
Ha/Wa	1.47 \pm 0.02 (1.19–1.82)	1.44 \pm 0.02 (1.16–1.93)	0.85	0.255
SH/BWH*	1.52 \pm 0.06 (0.87–2.37)	1.80 \pm 0.04 (1.36–2.89)	12.14	<0.001
PWH/BWH	0.25 \pm 0.01 (0.16–0.38)	0.25 \pm 0.00 (0.17–0.32)	0.62	0.731
BWH/W*	0.89 \pm 0.01 (0.77–1.09)	0.83 \pm 0.01 (0.73–0.98)	0.15	<0.001

Geometric morphometric analysis revealed a clear size and shape distinction between the two *Dianella* species. Shells of *D. thiesseana* were significantly larger (Centroid Size) than those of *D. schlickumi* ($F_{1,22} = 38.01$, $p < 0.0001$, Figure 8). The landmark-based analyses of individual shells also confirmed a shape distinction between the two species.

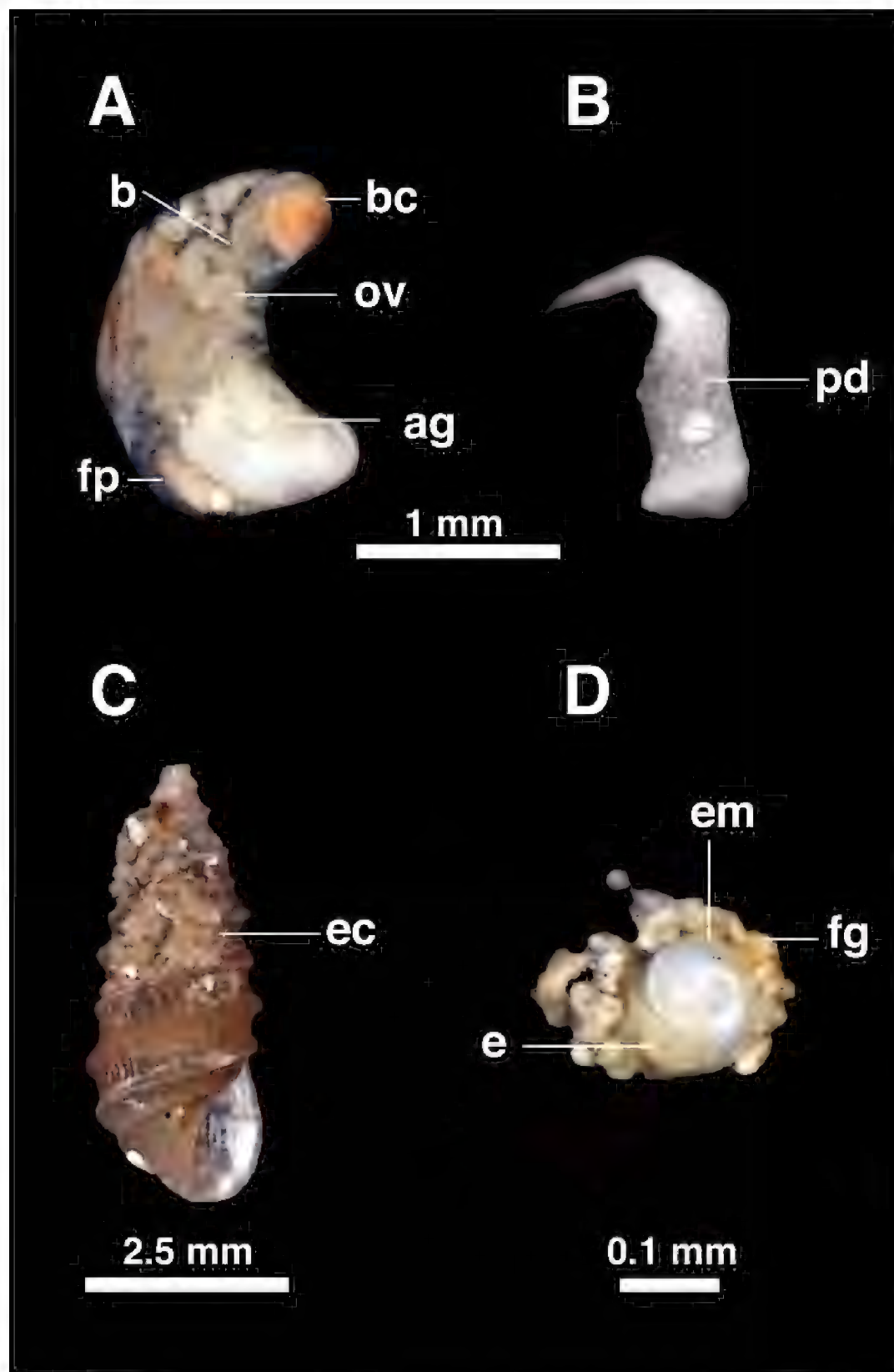


Figure 6. Genital anatomy and egg capsules of *Dianella schlickumi*. **A** Female genitalia **B** Penis **C** Shell bearing egg capsules covered by fine-grained material **D** Egg capsule with embryo. Abbreviations: ag—albumen gland; b—beak-shape dilatation of the oviduct; bc—bursa copulatrix; fp—faecal pellets; fg—fine-grained material; e—eye; ec—egg capsules; em—embryo; ov: renal oviduct; pd—penial duct.

ANOVA of Procrustes coordinates also showed a clear shape difference between the two species ($F_{32, 704} = 20.78, p < 0.0001$). PCA of Procrustes coordinates, which aimed at finding linear combinations maximising the total variance, revealed three principal components (PC1-PC3) explaining 88.1% of the total variance. The distinction between the two species was quite evident along the PC1 axis (Figure 9). In the CVA, which maximised differentiation amongst species (or predefined groups), the first and only extracted canonical variate

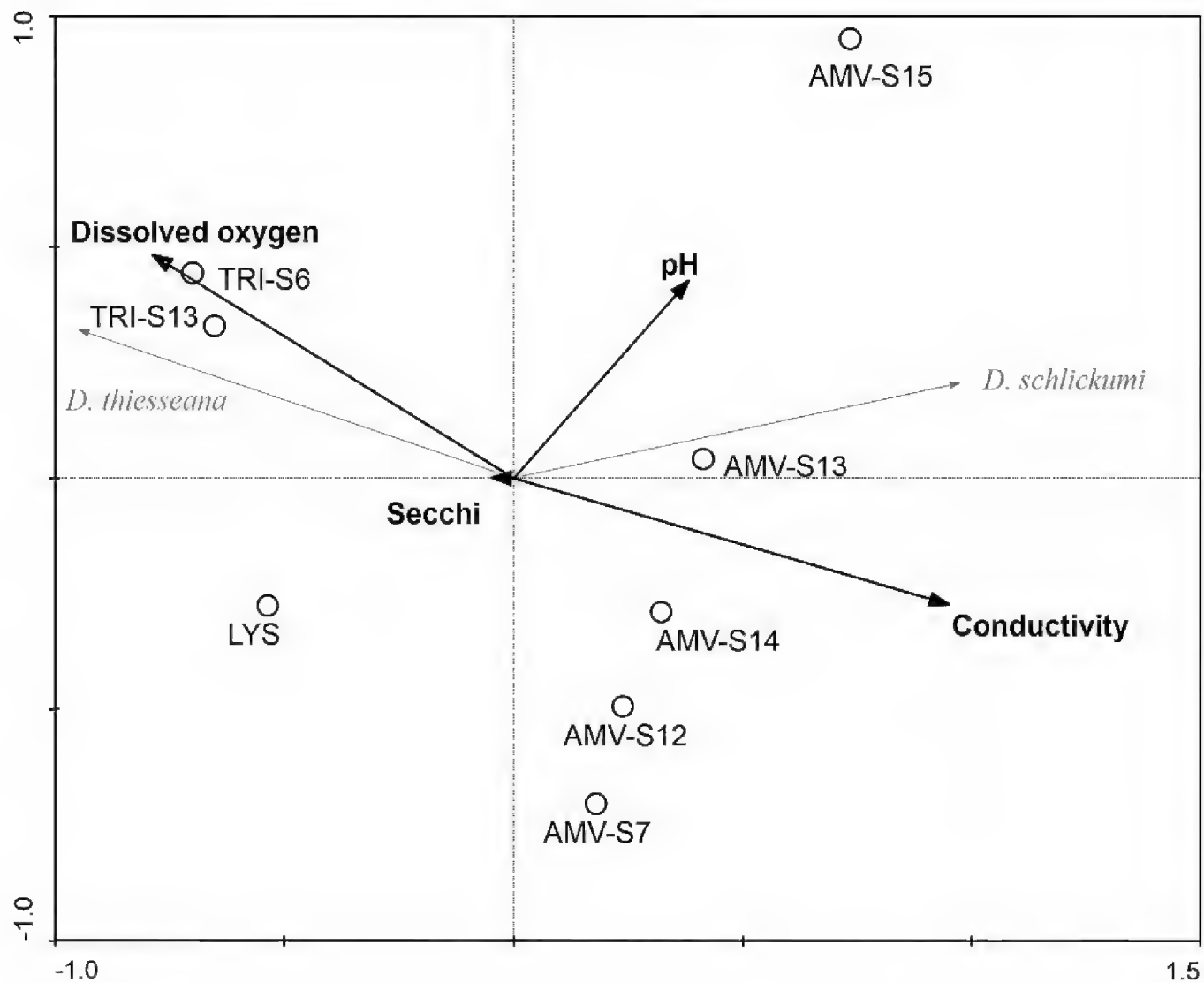


Figure 7. RDA ordination diagram of *Dianella thiesseana* and *Dianella schlickumi* abundances at sampling sites (circles) in relation to environmental parameters in lakes Trichonis (TRI), Lysimachia (LYS) and Amvrakia (AMV) in spring 2014 and 2015.

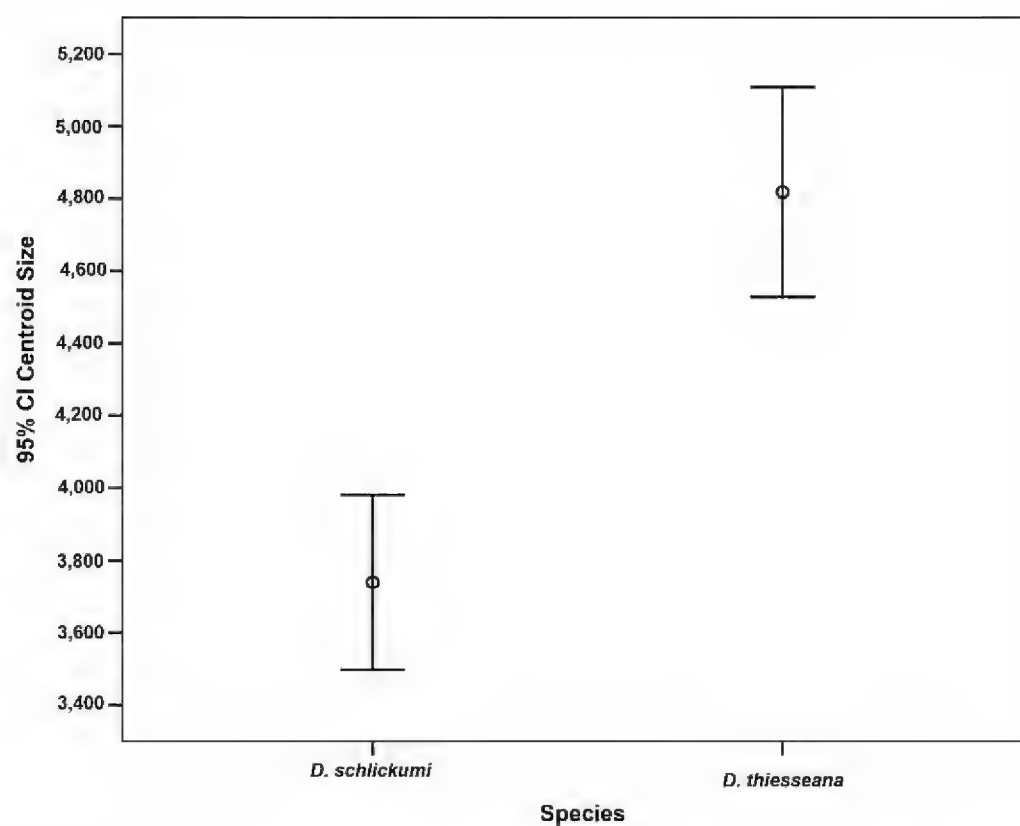


Figure 8. Means \pm 95% Confidence Intervals of centroid size for the two *Dianella* species and their ordinations based on landmark data.

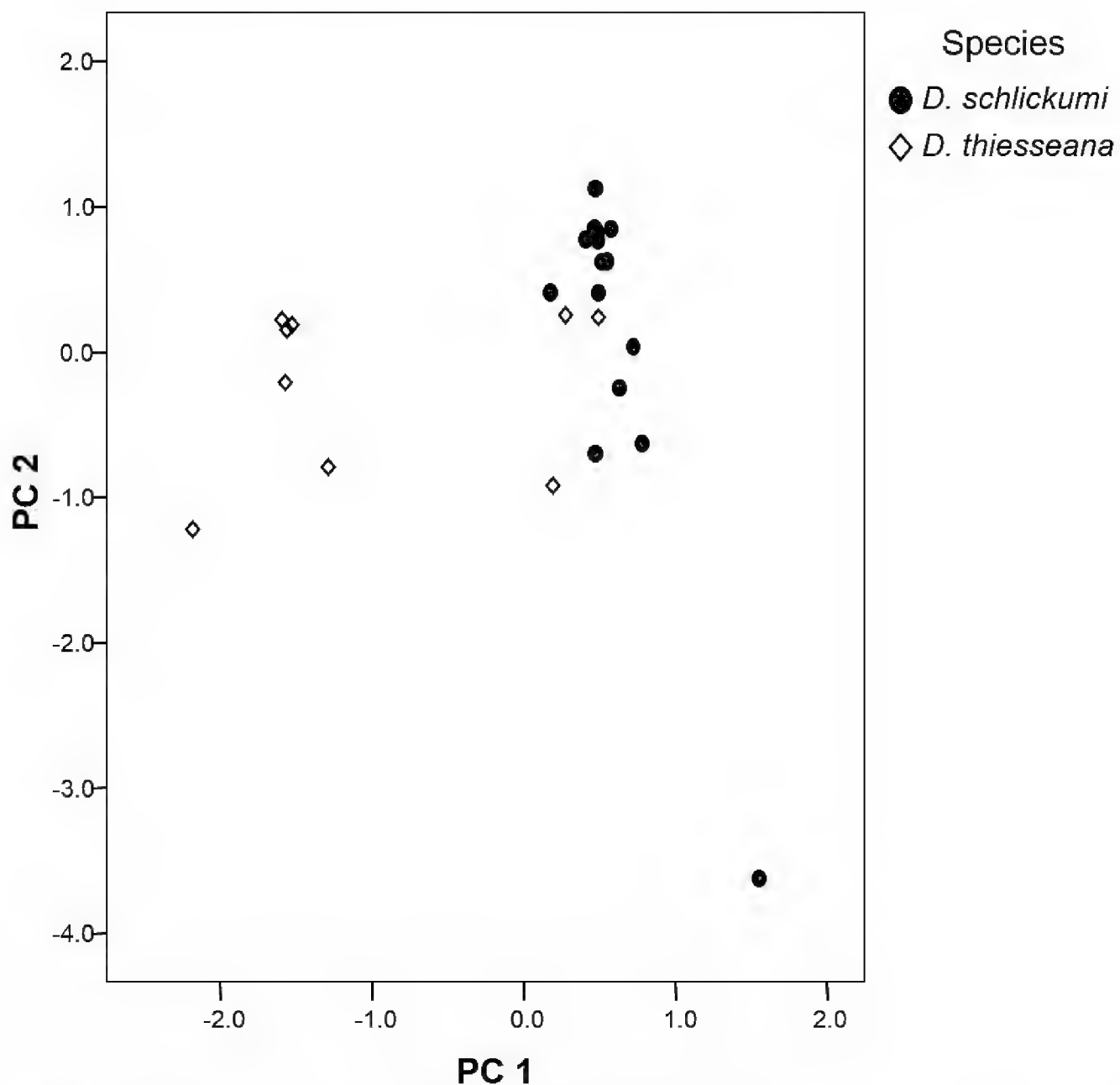


Figure 9. Principal component analysis of *Procrustes coordinates* for *Dianella schlickumi* (black circles) and for *Dianella thiesseana* (open diamonds).

(CV1) explained 100% of total variation and showed a much clearer separation than PCA between the two morphotypes along the CV axis (Figure 10). Clear shape discrimination was also found between *D. thiesseana* and *D. schlickumi* shells after a permutation Discriminant Function Analysis (T-squared = 236.3713, p (parametric) = 0.8335, p (permutation) < 0.0001). However, after a permutation test with 1,000 replicates, only 54.2% of “unknown” individuals (leave-one-out cross-validation) were correctly classified. Shape changes along the CV1 are shown in Figure 11 and portray the tendency for *D. thiesseana* to be slimmer compared to *D. schlickumi* and also to have a larger, i.e. elongated, aperture.

Environmental parameters

The Monte Carlo test in RDA analysis revealed two out of the six environmental parameters examined as significantly different between the two species ($p < 0.05$): conductivity and Secchi depth. Depth and water temperature were excluded from the analysis

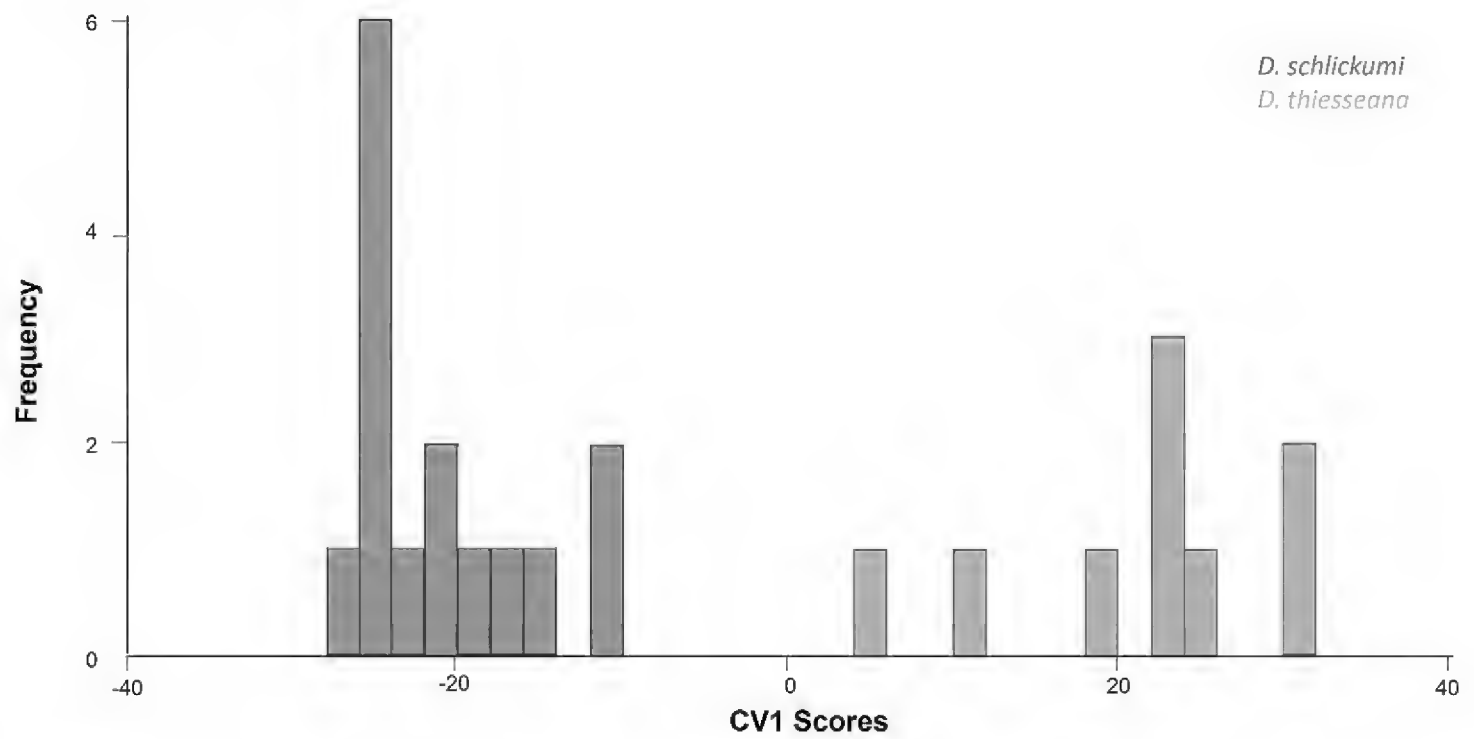


Figure 10. Canonical Variate Analysis: Dark-grey bins stand for *Dianella schlickumi* specimens and light-grey bins for *Dianella thiesseana* specimens.

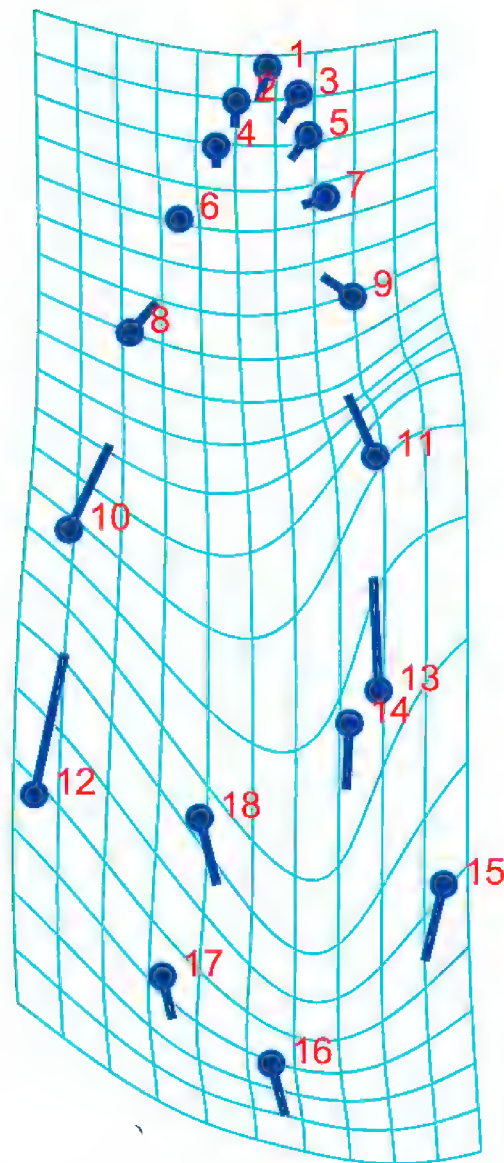


Figure 11. Shape changes along the CV1: the shifts of landmark positions are indicated by straight lines. Each line starts with a dot at the location of the landmark in the starting shape (i.e. *Dianella schlickumi*). The length and direction of the line indicate the movement of the respective landmark towards the final shape (i.e. *Dianella thiesseana*).

due to their high inflation factor (> 20). The first and all canonical axes were statistically significant ($p = 0.044$ and $p = 0.002$ respectively). The first two ordination axes of RDA explained 99.5% of the total species variance (Monte Carlo test, $p < 0.05$). Axis I (eigenvalue 0.929, $p < 0.05$) was related to conductivity and DO (intra-set correlation values 0.952 and -0.788 respectively) and Axis II (eigenvalue 0.066) to pH (intra-set correlation value 0.428). *D. schlickumi* specimens from Lake Amvrakia were ordered at the positive side of Axis I, having the highest conductivity values while *D. thiesseana* specimens from the other lakes were ordered at the negative side of Axis I and correlated with lower values of conductivity and higher values of DO (Figure 7, Table 2).

Discussion

Dianella schlickumi was recorded in the sublittoral zone of Lake Amvrakia in 1962 (Schütt 1962), but no exact details about this location were provided. The latest record confirming the presence of the species in the lake was before 1983 (Radoman 1983). After that, *D. schlickumi* was assumed to be extinct, as reported by Reischütz and Reischütz (2002), Szarowska et al. (2005) and Albrecht et al. (2006), based on sampling efforts in 1983, 2002 and 2003 respectively. Albrecht et al. (2011a) mentioned that the originally recorded area of *D. schlickumi* is the northern extent of Lake Amvrakia. At this part of the lake water, level fluctuations were high during the last years, due to insufficient connectivity with the main body of the lake. In fact, during extremely dry summers, this area is completely dry or water remains only in small pools. In such cases, the predation pressure (mainly from fish) is intense, resulting in local extinction of molluscs (Nilsson et al. 2008, Downing et al. 2010). Such rough conditions have probably negatively affected the population size and the distribution of *D. schlickumi* in this lake. Nevertheless, in our study, we were able to collect specimens of various ages at several sampling sites, indicating that this snail is still extant and thriving in its type locality.

Under the IUCN Red List guidelines (IUCN 2017), *D. schlickumi* is considered as “Critically Endangered (Possibly Extinct)” [following the criteria B1ab (i, iii)] (Albrecht et al. 2011a). However, since a viable population of this species was demonstrated in the present study, the tag “Possibly Extinct” should be removed from its assessment. The rediscovery of the species does not however improve its conservation status, as its extent of occurrence is estimated to be less than 100 km², restricted to the sublittoral zone of Lake Amvrakia (<10 km²). Moreover, *D. schlickumi* is known to exist only in a single location, where habitat degradation was also noticed due to agricultural practices and pesticide contamination (Thomatou et al. 2013 a, b). Consequently, the classification of Critically Endangered should be maintained, as the IUCN criteria continue to be met. As for *D. thiesseana*, it should also continue to be considered as Critically Endangered [B1ab (ii, iii) and 2ab (ii, iii)] since it has an extent of occurrence of 94 km² and an area of occupancy of 2 km². Its habitat is also degraded because of intensive agricultural practices, urban sewage, stock grazing land and small industries (Bertahas et al. 2006).

Both species thrive on soft substrate (Albrecht et al. 2011a, b). *D. schlickumi* seemed to prefer slightly shallower habitats, from 3 m to 15 m, while *D. thiesseana* had a range of distribution from 2.5 m to 27 m. The species absence from autumn surveys may be due to depth migrations often observed in gastropods (Brown 2001). Even though this study has confirmed the rediscovery of *D. schlickumi*, it does not clarify in detail the accurate distribution of the species, which might be the subject of future research.

The anatomical characters of *D. schlickumi* known from literature (Schütt 1962) (i.e. the radula) and those described in the present paper (i.e. the ctenidium and osphradium, the central nervous system, the gastric caecum, the rectum and the male and female reproductive organs) are similar to those reported by Radoman (1983) in the diagnosis of the family Pyrgulidae Brusina 1881 (nominal subfamily Pyrgulinae) and of the genus *Dianella* Gude, 1913 [(type species *Dianella thiesseana* (Kobelt, 1878)]. Similarities have also been found between *D. schlickumi* and *D. thiesseana* in the nervous system, the gastric caecum and the female reproductive organs as they are described for the latter species by Szarowska et al. (2005, pp. 22–31 and related figures). However, a notable difference exists in the shape of the penis between the two species: the penis of *D. thiesseana* is wide, with a triangular shape due to its expanded base and has a clear outgrowth on its left side (Szarowska et al. 2005, p. 29, figs 32–37). The penis of *D. schlickumi* does not have an expanded base and is gradually tapering without any distinct outgrowth.

D. schlickumi and *D. thiesseana* are discriminated due to their shell dimensions; the first one is smaller than the second (Radoman 1983). In our study, traditional and geometric morphometric analyses support the assumption that *D. thiesseana* and *D. schlickumi* are quite distinguishable in terms of external morphology. However, neither in the original description of the two taxa (Kobelt 1878 and Schütt 1962 respectively) nor in the subsequent studies (Radoman 1983, Szarowska et al. 2005), shell shape appears to be significantly different between the two taxa. Moreover, Szarowska et al. (2005) reported that the shell characters and dimensions of *D. schlickumi*, described in the literature and those of *D. thiesseana* from Lake Lysimachia, lie within the range observed for *D. thiesseana* from Lake Trichonis. Even though, the intra-subspecific morphometric variation is quite high, the two species are actually distinct in size and shape (Figures 8–10). Therefore, in the “*Dianella*” case, geometric morphometric analysis has been proved to be a useful tool for exploring patterns of species diversity and seems promising for examining morphological adaptations of these species. Specifically, it is clear that the Centroid Size, derived from Geometric Morphometric Analysis, can be used safely to distinguish those two species.

In some cases, the environmental parameters set limits (Pyron and Brown 2015) and may influence (Økland 1983, Pip 1986, Falniowski 1987) the distribution of freshwater snails. The analysis performed revealed conductivity, DO and pH as the most significant parameters explaining the current species variance. *D. thiesseana*, according to RDA, seems to prefer more oxygenated waters with lower conductivity values and pH than *D. schlickumi*. Conductivity, dissolved oxygen (e.g. Çabuk et al. 2004, Kazibwe et al. 2006, Cloherty and Rachlin 2011) and pH (Økland 1983, Pip 1986, Çabuk et al. 2004) are also referenced as key environmental proxies explaining

gastropods distribution in water bodies. Biological interactions and/or historical factors, which need further surveys and analyses, may also be responsible.

Both *Dianella* species require urgent conservation measures to be enforced and suitable management plans to be implemented in the whole studied area, focusing on the protection of these species and the improvement of their habitats. The studied lakes form an ecologically important complex as Special Conservation areas (Council Directive 92/43/EEC), listed in the NATURA 2000 network. However, the integrity of local mollusc populations in these ecosystems is threatened due to the presence of invasives; *Ferrissia fragilis* (Tryon, 1863) and *Physa acuta* Draparnaud, 1805 in lakes Trichonis and Lysimachia (Albrecht et al. 2014) and of *Potamopyrgus antipodarum* (J.E. Gray, 1843) in Lake Trichonis (Radea et al. 2008). The invasive impacts upon native molluscs may be direct, through competition for food and space (Carlsson et al. 2004) or indirect, through changes in ecosystem function and/or parasitism (Brown et al. 2008). The antagonistic behaviour of *P. antipodarum* and *P. acuta* is already well-known (e.g. Schreiber et al. 2003, Zukowski and Walker 2009), although no such outcomes have been documented for *F. fragilis* until now (Albrecht et al. 2014). Available knowledge on the effects of invasive species on native mollusc species and especially on *D. thiesseana*, is insufficient to estimate the possible threats.

One of the main pressures identified in lakes Amvrakia, Trichonis and Lysimachia is eutrophication from agriculture run-off and wastewaters (Bertahas et al. 2006, Thomatou et al. 2013 a, b). *Dianella* inhabits the sublittoral zone of these lakes, which is mainly affected by eutrophication (Pilotto et al. 2012). The development of water quality indices based on zoobenthic communities and related to eutrophication will help to assess the ecological status of these water bodies and prevent their further deterioration. The impacts could be minimised by preserving natural vegetation or establishing buffer zones. Only low-impact activities should be allowed near the lakes and harmful practices should be prohibited or regulated to a more distant zone. Furthermore, it is crucial to restore the type locality of *D. schlickumi* by maintaining the connectivity of the northern extension of Lake Amvrakia with the main lake water body, thus preventing the species' historical distribution area to become dry. Additional measures to reduce water abstraction could help the maintenance of the ecological water level in the lakes, securing slight seasonal fluctuations and preventing aquatic biodiversity in general (Zohary and Ostrovsky 2011). Further temporal and spatial surveys which will identify the ecological features, the ecosystem functions and the habitat preferences of the studied species could significantly support conservation efforts. Environmental education and citizen awareness could additionally support conservation efforts for both endemic gastropods.

Conclusion

Our results confirm the rediscovery of the narrow endemic species *Dianella schlickumi* in Lake Amvrakia since past sampling efforts did not detect it for more than

30 years. Moreover, they provide basic knowledge on its anatomical, morphological and distributional patterns and its discrimination from *Dianella thiesseana*. The two species are discriminated by their shell size and shape. Moreover, the presence of *Dianella thiesseana* is confirmed in lakes Trichonis and Lysimachia. We conclude that further conservation measures have to be implemented for effective protection of both species.

Acknowledgements

We would like to thank Mrs Efi Mavromati from the Greek Biotope/Wetland Centre for providing us with data from the National Greek Monitoring Programme for the presence of *Dianella thiesseana* in Lake Lysimachia.

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